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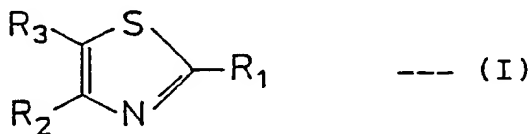
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D-8500 Nürnberg(DE)(54) **Thiazole derivatives.**(57) A thiazole derivative represented by the formula
I, pharmaceutically acceptable acid addition salt

thereof, and a process for preparation thereof



wherein R_1 represents $-\text{COOR}_4$, $-\text{CON} \begin{smallmatrix} R_5 \\ R_6 \end{smallmatrix}$ or cyano

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wherein R_4 represents hydrogen or lower alkyl, and

R_5 and R_6 may be same or different and represent hydrogen, lower alkyl, aryl, amino-lower alkyl, N-lower alkylamino-lower alkyl, N,N-di-lower alkylamino-lower alkyl or R_5 and R_6 are joined to form N-containing heterocyclic group;

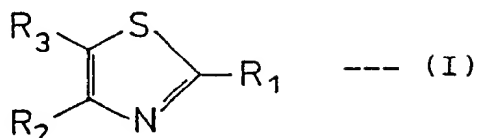
R_2 represents hydrogen or lower alkyl; and

R_3 represents N-containing heterocyclic group.

THIAZOLE DERIVATIVES

BACKGROUND OF THE INVENTION

The present invention relates to a thiazole derivative represented by the formula I, pharmaceutically acceptable acid addition salt thereof, and a process for preparation thereof



wherein R_1 represents $-\text{COOR}_4$, $-\text{CON} \begin{smallmatrix} R_5 \\ R_6 \end{smallmatrix}$ or cyano

wherein R_4 represents hydrogen or lower alkyl, and

R_5 and R_6 may be same or different and represent hydrogen, lower alkyl, aryl, amino-lower alkyl, N-lower alkylamino-lower alkyl, N,N-di-lower alkylamino-lower alkyl or R_5 and R_6 are joined to form N-containing heterocyclic group;

R_2 represents hydrogen or lower alkyl; and

R_3 represents N-containing heterocyclic group.

Some thiazole derivatives having a cardiotoxic activity are known as described in Japanese Patent Applications laid open under Nos. 134417/1982, 16889/1984 and 193878/1984 and so on. Meanwhile, thiazole derivatives in general are prepared by a method as described in Japanese Patent Applications laid open under Nos. 34241/1982, 49969/1974 and so on. That is, a thioamide is reacted with an α -halocarbonyl compound to form a compound of thiazole derivative. The known thiazole derivatives having a cardiotoxic activity is prepared by this well known method.

However, there is no teaching in the prior art of any cardiotoxically active compounds having any chemical structure comparable to or suggestive of instantly claimed compound.

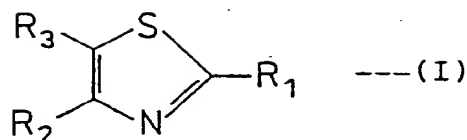
Further, such known technique is unsuitable for synthesizing the compound of the formula I.

SUMMARY OF THE INVENTION

As the result of researches, we, the inventors succeeded in synthesizing new thiazole derivatives which are useful as cardiotoxic agents. More specifically, one aspect of this invention is to provide thiazole derivatives having carbonyl, unsubstituted or substituted aminocarbonyl or cyano at position 2 and having nitrogen-containing heterocyclic group at position 6 and and pharmaceutically acceptable acid addition salt thereof. Another aspect of this invention is to provide a cardiotoxic agent comprising said thiazole derivatives and pharmaceutically acceptable acid addition salt thereof as effective component. A further aspect of this invention is to provide a process for preparation of said thiazole derivatives and pharmaceutically acceptable acid addition salt thereof.

DETAILED DESCRIPTION

The thiazole derivative of the present invention is represented by the following formula I



wherein R_1 represents $-\text{COOR}_4$, $-\text{CON}(\text{R}_5)(\text{R}_6)$ or cyano

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wherein R_4 represents hydrogen or lower alkyl, and R_5 and R_6 may be same or different and represent hydrogen, lower alkyl, aryl, amino-lower alkyl, N-lower alkylamino-lower alkyl, N,N-di-lower alkylamino-lower alkyl or R_5 and R_6 are joined to form N-containing heterocyclic group;

R_2 represents hydrogen or lower alkyl; and

R_3 represents N-containing heterocyclic group.

The term "lower" as used herein refers to from 1 to 6 carbon atoms unless otherwise indicated.

The term "lower alkyl" as used herein means alkyl group having from 1 to 6 carbon atoms which may be arranged as straight or branched chains. The "lower alkyl" may be methyl, ethyl, n-propyl, iso-propyl, n-butyl, iso-butyl, tert-butyl, n-pentyl, n-hexyl or the like.

The term "aryl" as used herein means aryl group which may be unsubstituted or may bear one or two substituents selected from lower alkyl, lower alkoxy (e.g. methoxy, ethoxy), halogeno (e.g. chloro, bromo), cyano, nitro, hydroxy and so on. The "aryl" may be phenyl, tolyl, xylyl, mesityl, cumenyl, biphenyl, hydroxyphenyl or the like.

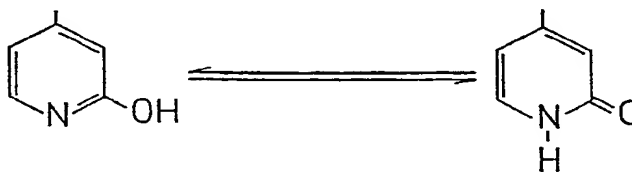
The term "amino-lower alkyl" as used herein means lower alkyl group substituted with one or two of amino groups. The "amino-lower alkyl" may be aminomethyl, diaminomethyl, aminoethyl, diaminoethyl, amino-n-hexyl or the like.

The term "N-lower alkylamino-lower alkyl" as used herein means amino-lower alkyl group substituted with one lower alkyl group. The "N-lower alkylamino-lower alkyl" may be N-methylaminomethyl, N-methylaminoethyl, N-ethylaminoethyl, N-n-hexylaminoethyl or the like.

The term "N,N-di-lower alkylamino-lower alkyl" as used herein means amino lower alkyl group substituted with two lower alkyl groups. The "N,N-di lower-alkyl amino-lower alkyl" may be N,N-dimethylaminomethyl, N,N-dimethylaminoethyl, N-methyl-N-n-hexylaminomethyl or the like.

The term "N-containing heterocyclic group" as used herein means nitrogen-containing, 5 or 6 membered mono or condensed heterocyclic ring group which may be unsubstituted or may bear one or two substituents selected from lower alkyl, lower alkoxy, halogeno, cyano, nitro, aldehyde, acyl (e.g. formyl, acetyl), hydroxy and so on. The "N-containing heterocyclic group" may be pyridinyl, pyridazinyl, pyrimidinyl, pyrazinyl, quinolinyl, isoquinolinyl, cinnolinyl, quinazolinyl, quinoxalinyl, phthalazinyl, acridinyl, methylpyridinyl, methoxypyridinyl, chloropyridinyl, formylpyridinyl, cyanopyridinyl, nitropyridinyl, acetylpyridinyl, hydroxypyridinyl, methoxyquinolinyl, methylpyridazinyl, piperidino, morpholino or the like.

As to the formula I, it is to be noted that some of the compounds may be alternatively represented by its tautomers, e.g. "2-hydroxypyridinyl" and "2-oxo-pyridinyl". Both of said moieties are in the state of tautomeric equilibrium as represented by the following equilibriums.



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Accordingly it is to be understood that both of such isomers are included within the same category of the formula I.

In the present specification, the formula I includes the group of such tautomeric isomers which is however represented by one of the expressions.

The compounds according to the present invention are for example as follows:

. methyl 4-methyl-5-(4-pyridinyl)-thiazole-2-carboxylate;

. 4-methyl-5-(4-pyridinyl)-thiazole-2-carboxamide;

. N,4-dimethyl-5-(4-pyridinyl)-thiazole-2-carboxamide;

. 4-methyl-5-(4-pyridinyl)-thiazole-2-carboxanilide;

. 4-methyl-N-aminomethyl-5-(4-pyridinyl)-thiazole-2-carboxamide;

. 4-methyl-N-(2-methylaminoethyl)-5-(4-pyridinyl)-thiazole-2-carboxamide;

. 4-methyl-N-(2-dimethylaminoethyl)-5-(4-pyridinyl)-thiazole-2-carboxamide;

. N,N-diethyl-4-methyl-5-(4-pyridinyl)-thiazole-2-carboxamide;

. 2',6'-dimethyl-4-methyl-5-(4-pyridinyl)-thiazole-2-carboxamide;

. 1-[4-methyl-5-(4-pyridinyl)-2-thiazolylcarbonyl]-piperidine;

. 2-cyano-4-methyl-5-(4-piperidinyl)-thiazole;

. methyl 4-methyl-5-(4-pyridazinyl)-thiazole-2-carboxylate;

. 4-methyl-5-(4-pyridazinyl)-thiazole-2-carboxamide;

. 1-[4-methyl-5-(4-pyridazinyl)-2-thiazolylcarbonyl]-piperazine;

. 2-cyano-4-methyl-5-(4-pyridazinyl)-thiazole;

. methyl 4-methyl-5-(4-quinoliny)-thiazole-2-carboxylate;

. 4-methyl-5-(4-quinoliny)-thiazole-2-carboxamide;

. 2-cyano-4-methyl-5-(4-quinoliny)-thiazole;

. methyl 4-methyl-5-(3-methyl-4-pyridinyl)-thiazole-2-carboxylate;

5 . 4-methyl-5-(3-methyl-4-pyridinyl)-thiazole-2-carboxamide;

. 2-cyano-4-methyl-5-(3-methyl-4-pyridinyl)-thiazole;

10 . methyl 4-methyl-5-(3-acetyl-4-pyridinyl)-thiazole-2-carboxylate;

. 4-methyl-5-(3-acetyl-4-pyridinyl)-thiazole-2-carboxamide;

. 2-cyano-4-methyl-5-(3-acetyl-4-pyridinyl)-thiazole;

. methyl 4-methyl-5-(3-chloro-4-pyridinyl)-thiazole-2-carboxylate;

. 4-methyl-5-(3-chloro-4-pyridinyl)-thiazole-2-carboxamide;

. 2-cyano-4-methyl-5-(3-chloro-4-pyridinyl)-thiazole;

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. methyl 4-methyl-5-(3-formyl-4-pyridinyl)-2-carboxylate;

. 4-methyl-5-(3-formyl-4-pyridinyl)-thiazole-2-carboxamide;

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. 2-cyano-4-methyl-5-(3-formyl-4-pyridinyl)-thiazole;

. methyl 4-methyl-5-(3-cyano-4-pyridinyl)-thiazole-2-carboxylate;

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. 4-methyl-5-(3-cyano-4-pyridinyl)-thiazole-2-carboxamide;

. 2-cyano-4-methyl-5-(3-cyano-4-pyridinyl)-thiazole;

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. methyl 4-methyl-5-(6-methoxy-4-quinoliny)-thiazole-2-carboxylate;

. 4-methyl-5-(6-methoxy-4-quinoliny)-thiazole-2-carboxamide;

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. 2-cyano-4-methyl-5-(6-methoxy-4-quinoliny)-thiazole;

. methyl 4-methyl-5-(2-thiazolyl)-thiazole-2-carbox-

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ylate;

. 4-methyl-5-(2-thiazolyl)-thiazole-2-carboxamide;

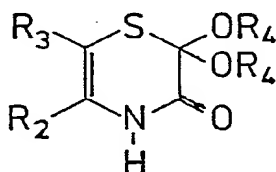
. 2-cyano-4-methyl-5-(2-thiazolyl)-thiazole;

and the like.

The thiazole derivative of the formula I is prepared according to the following process.

As to the following process, the pressure is under atmospheric pressure unless otherwise indicated.

First of all, the 2,2-di-lower alkoxy-1,4-thiazine derivative of the formula II is prepared

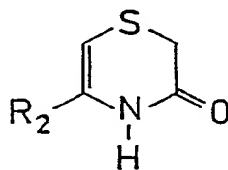


--- (II)

wherein R₂, R₃ and R₄ are as defined above.

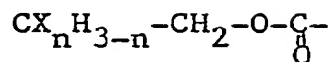
The compound of the formula II is prepared according to the following process.

When a known 1,4-thiazine derivative of the formula



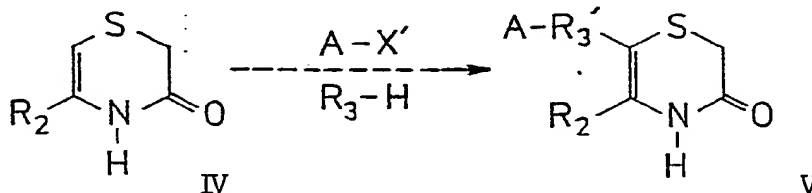
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wherein R₂ is as defined above, is reacted with a known compound of the formula A-X' (in which A represents a residue



in which X represents a halogen atom and n is a number of 0 to 3, and X' represents a halogen atom which may be the same as or different from

X) and a known compound of the formula R₃-H (in which R₃ is as defined above), a 1,4-thiazine derivative of the formula V is obtained.



wherein R_2 , R_3 and A are as defined above and R_3' represents dihydro form residue of N-containing heterocyclic group.

This reaction is completed by merely stirring the compound of the formula VI, more than equal mole concentration of the compound A-X' and more than equal mole concentration of the compound of the formula R_3 -H at room temperature for 1 to 7 hours.

The solvent, which is used in this reaction, may be nitrile such as acetonitrile, ether such as tetrahydrofuran, or halogenated hydrocarbon such as dichloromethane or the like. Incidentally, the compound having N-containing heterocyclic group may also be used as a solvent.

The compound having N-containing heterocyclic group may be unsubstituted or may bear one or two substituents selected from lower alkyl, lower alkoxy, halogeno, cyano, nitro, al-

dehyde, acyl, hydroxy, alkoxycarbonyl and so on. The compound may be pyridine, pyridazine, pyrimidine, pyrazine, quinoline, isoquinoline, cinoline, quinazoline, quinoxaline, phthalazine, acridine, phenazine, piperidine, piperazine, morpholine, 3-methoxypyridine, 3-chloropyridine, pyridine-3-aldehyde, nicotinonitrile, 3-acetylpyridine, methyl nicotinate, 3-nitropyridine, 3-hydroxypyridine, 6-methoxyquinoline, 3-methylpyridazine or the like.

Then the compound of the formula IV is obtained by aromatization of the compound of the formula V. This may be effected in various processes; a preferred process is as follows.

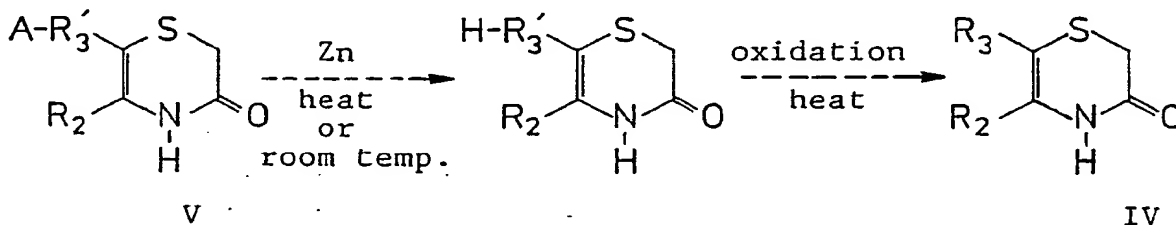
The compound of the formula V is reacted with sulfur at elevated temperature to obtain the 1,4-thiazine derivative of the formula IV.



wherein R_2 , R_3 , A and R_3' are as defined above.

This reaction proceeds well by stirring the compound of the formula V with from half to 5 fold amount of sulfur and heating the mixture at 120 to 200°C for 0.5 to 8 hours. Generally, no solvent is needed in this reaction; however, N,N-dimethylformamide, dimethyl sulfoxide, xylene, (o-, m-, p-) dichlorobenzenes, diglyme, etc. may be employed.

Alternatively, the compound of the formula IV may be prepared according to a process that the compound of the formula V is reacted with zinc at room temperature or elevated temperature. Some of the compound of the formula V is reacted with zinc to obtain thiazole derivative having dihydro form residue of N-containing heterocyclic group at position 6. In this case, it is necessary to oxidize a reaction intermediate by an oxidizing agent. This process is represented by the following formula



wherein R_2 , R_3 , A and R_3' are as defined above.

This reaction proceeds well by stirring the compound of the formula V with excessive amount of zinc and stirring the mixture at room temperature or elevated temperature up to 80°C for 1 to 4

hours. A solvent used in this reaction may be an organic solvent in the form of mixture with water; the organic solvent may be carboxylic acid such as formic acid, nitrile such as acetonitrile, ether such as tetrahydrofuran, sulfoxide such as dimethylsulfox-

- ide or amide such as methylformamide or the like. The oxidizing agent used in this reaction may be 2,3-dichloro-5,6-dicyanobenzoquinone (DDQ) or the like.

Purification of the compounds of the formula V and VI may be accomplished by recrystallization from lower alcohol such as methanol, ethanol or isopropanol, ketone such as acetone, halogenated hydrocarbon such as chloroform, carboxylic acid ester such as ethyl acetate, aromatic hydrocarbon such as benzene, ether such as diethylether or nitrile such as acetonitrile or the like.

Alternatively, such purification may also be accomplished by column chromatography or thin layer chromatography. In this operation, silica gel having particles size of 100-200 mesh such as Wakogel C-200 (manufactured by Wako Junyaku Kabushiki Kaisha in Japan), silica gel having particle's average porous diameter of 40-63 μm such as silica gel 60 Lichloprep 60 (manufactured by

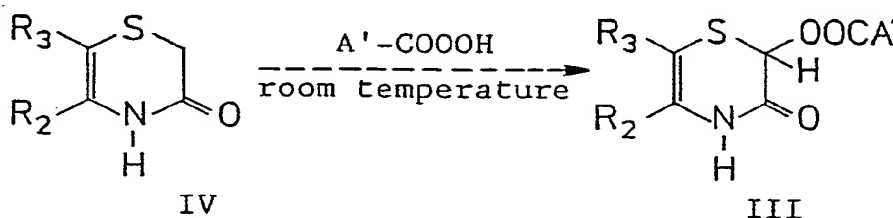
Merck & Co. Inc. in USA) or fluoresces a light blue color in the region of 254 nm such as Merck TCL plate silica gel 60F₂₅₄ (manufactured by Merck & Co. Inc. in USA) is preferable to use.

Incidentally, the compound of the formula V may be directly used in the subsequent reaction without purification.

The known starting material 1,4-thiazine derivative of the formula VI may be prepared according to the processes proposed by H. Sokol et al. in J. Am. Chem. Soc., 70, 3517 (1948), C. R. Johnson et al. in J. Heterocycl. Chem., 6, 247-249 (1969), and G. V. Rao et al. in Synthesis, 136 (1972).

Then, the thiazine derivative having substituent group such as lower alkoxy, halogeno, acyloxy or hydroxy or the like at position 2 may be prepared according to the process by M. Hojo et al. in Synthesis 312, 424 (1982).

Thus, the compound of the formula IV is reacted with the compound of the formula A'-COOOH - (in which A' represents lower alkyl, alicyclic residue or aryl), i.e. peroxycarboxylic acid to obtain the compound of the formula III.



wherein R₂, R₁ and A' are as defined above.

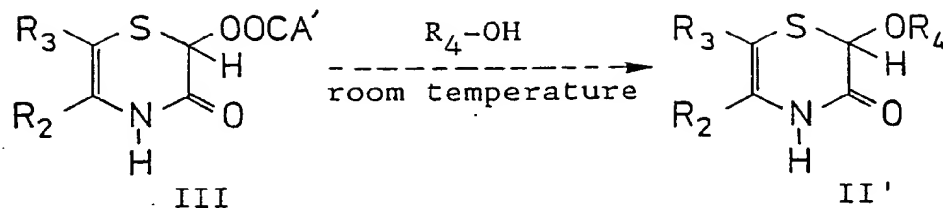
This reaction is completed by merely stirring the compound of the formula IV and more than equal mole concentration of the compound of the formula A'-COOOH under ice-cooling for a few minutes.

The solvent, which is used in this reaction, may be halogenated hydrocarbon such as dichloromethane, nitrile such as acetonitrile, ether such as tetrahydrofuran, sulfoxide such as dimethylsulfoxide or amide such as dimethylformamide or the like.

As for the peroxycarboxylic acid, aliphatic peroxycarboxylic acid such as performic acid or peracetic acid, alicyclic peroxycarboxylic acid such as cyclohexaneperoxycarboxylic acid, or aromatic per-

oxycarboxylic acid such as perbenzoic acid or monoperoxyphthalic acid may be employed. The peroxycarboxylic acid having substituent such as lower alkyl, lower alkoxy, halogeno, cyano, nitro, aldehyde, acyl, hydroxy or the like may also be employed. In view of the state of the reaction system (the reactivity and degree of dissociation) and the easy availability, m-chloroperbenzoic acid is especially preferred.

Then, the compound of the formula III is reacted with the nucleophilic reagent of the formula R₄-OH (in which R₄ is as defined above) to obtain the compound of the formula II'.



wherein R₂, R₃, R₄ and A' are as defined above.

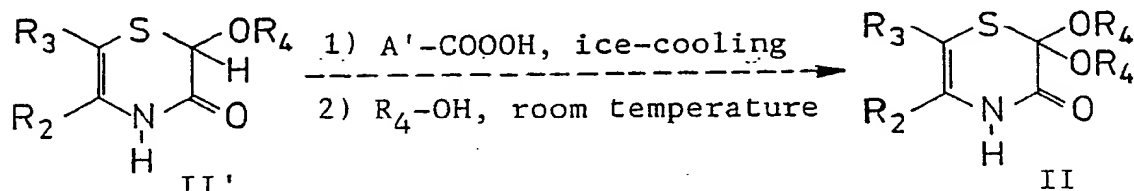
This reaction is completed by merely stirring the compound of the formula III and more than equal mole concentration of the compound of the formula R₄-H at room temperature for more than 1 day. Alternatively, the reaction may be carried out at 50-70°C.

In this reaction, the compound of the formula R₄-H may be used as a solvent

Incidentally, amide such as dimethylformamide, nitrile such as acetonitrile, ether such as tetrahydrofuran, sulfoxide such as dimethylsulfoxide or halogenated hydrocarbon such as dichloromethane or the like may be used as a solvent.

As for the compound of the formula R₄-OH, lower alcohol such as methanol, ethanol, isopropanol or the like may be employed.

Then, the compound of the formula II' is reacted with more than equal mole concentration of the formula A'-COOOH and more than two mole concentration of the formula R₄-OH to obtain the compound of the formula II.



wherein R₂, R₃, R₄ and A' are as defined above.

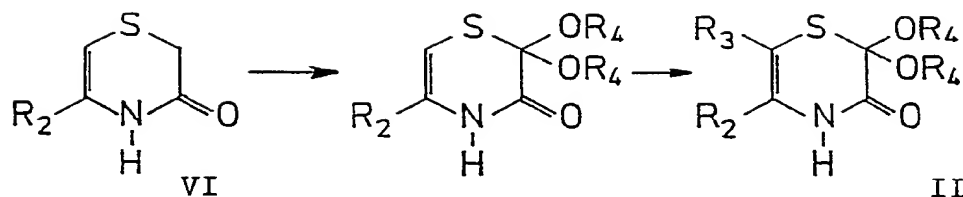
The compound of the formula III, the compound of the formula II' and the compound of the formula II may be used in the subsequent reaction without purification.

Incidentally, purification of these compounds may be accomplished by recrystallization from lower alcohol such as methanol, ethanol or isopropanol, halogenated hydrocarbon such as chloroform or carboxylic acid ester such as ethyl acetate or the like.

Alternatively, such purification may also be accomplished by alumina column chromatography. In this operation, alumina having 300 mesh is preferable to use. And, halogenated hydrocarbon such as chloroform or carboxylic acid ester such as ethyl acetate or the like may be used as an eluent.

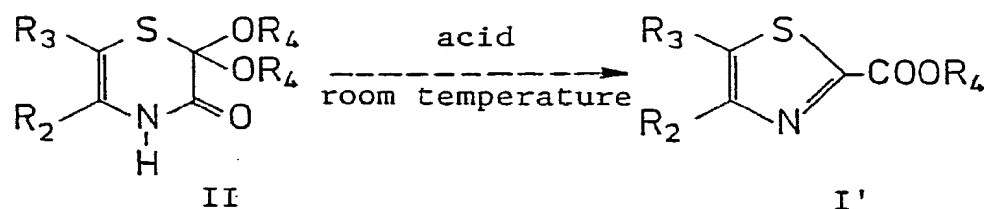
The thiazine derivative having substituent at position 2, i.e. the compound of the formula II, may be also obtained by the following process. One mole concentration of the compound of the formula IV is reacted with more than two mole concentration of the compound of the formula A'-COOOH and more than two mole concentration of the compound of R₄-OH in a similar manner as above.

Alternatively, the compound of the formula II may be obtained by introduction of -OR₄ group position at 2 in the same manner as above, then introduction of -R₂ group position at 6 in the same manner as above.



wherein, R_2 , R_3 , R_4 are as defined above.

The compound of the formula II is reacted with an acid to obtain the compound of the formula I' i.e. thiazole derivative having lower alkoxycarbonyl group at position 2.



wherein R_2 , R_3 and R_4 are as defined above.

This reaction is completed by merely allowing the compound of the formula II with more than equal mole concentration of the acid and the mixture to stand at room temperature for more than 10 minutes. The acid used in this reaction may be inorganic acid such as hydrochloric acid, sulfuric acid, etc. or organic acid such as acetic acid, etc. In view of the state of the reaction system (the reactivity and degree of dissociation) and the easy availability, hydrochloric acid is especially preferred.

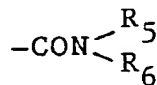
This reaction is proceeded by adding the compound of the formula II into the acid which is diluted with water.

The solvent used in this reaction is water.

Purification of the compound of the formula I' may be accomplished by recrystallization from aromatic hydrocarbon such as benzene or alicyclic hydrocarbon such as cyclopentane, cyclohexane or the like or a mixture of two or more of these solvents.

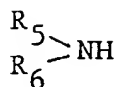
Alternatively, such purification may be accomplished by column chromatography or thin layer chromatography.

Optinally, the $-\text{COOR}_4$ group (in which R_4 is as defined above) of the compound of the formula I' may be converted into



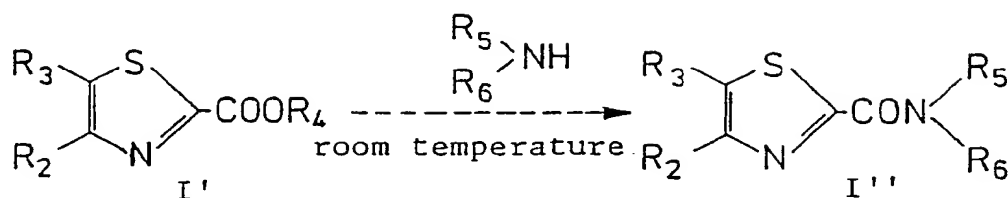
group (in which R_5 and R_6 are as defined above) or cyano group by methods known in the art. Thus the compound of the formula I'', i.e. thiazole derivative having unsubstituted or substituted aminocar-

bonyl group at position 2, is prepared by reacting the compound of the formula I' with the compound of the formula



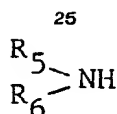
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(in which R_5 and R_6 are as defined above).



wherein R_2 , R_3 , R_4 , R_5 and R_6 are as defined above. 20

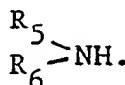
This reaction is completed by merely stirring the compound of the formula I' and more than equal mole concentration of the compound of the formula



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at room temperature for more than 1 hour, preferably about 10 hours. 30

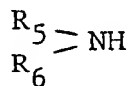
Alternatively, it may be completed by once converting the compound of the formula I' into a reaction intermediate having chlorocarbonyl group and then reacting the same with the compound of the formula 35



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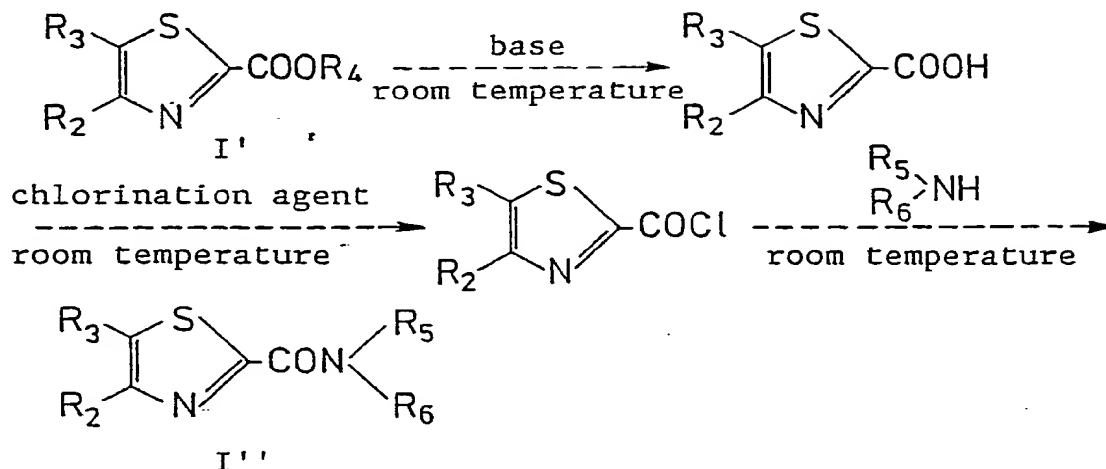
More specifically, first, the compound of the formula I' and a base such as sodium hydroxide aqueous solution or potassium hydroxide aqueous solution are stirred at room temperature to obtain a thiazole derivative having carboxy group at position 2. Secondly, the compound having carboxy group at position 2 and more than equal mole of chlorina-

tion agent such as thionyl chloride are stirred at room temperature for more than 10 hours to obtain a thiazole derivative having chlorocarbonyl group at position 2. Thirdly, the compound having chlorocarbonyl group at position 2 is reacted with more than equal mole of the compound of the formula 45



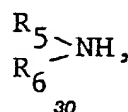
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at room temperature for more than 10 hours to obtain the compound of the formula I'. This process is represented by the following formula



wherein R₂, R₃, R₄, R₅ and R₆ are as defined above.

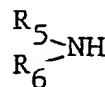
As for the compound represented by the formula



ammonia water, lower alkyl amine such as methylamine or ethylamine, aromatic amine such as aniline, toluidine or xyldine, di-amine such as N,N-dimethylaminoethylamine, secondary amine such as N,N-dimethylamine, N,N-diethylamine or heterocyclic compound having nitrogen atoms such as piperidine, piperazine or morpholine or the like

may be used in this reaction. As for the solvent, lower alcohol such as methanol or ethanol, ether such as tetrahydrofran, aromatic hydrocarbon such as toluene or xylene or water or the like may be used in this reaction.

Alternatively, the compound of the formula



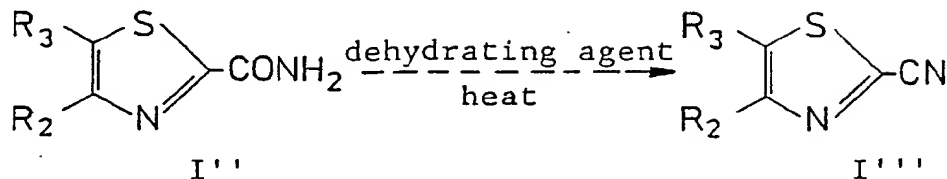
may be used as a solvent.

Purification of the compound of the formula I'' may be accomplished by recrystallization from lower alcohol such as methanol, ethanol or iso-propanol, aromatic hydrocarbon such as benzen, alicyclic hydrocarbon such as cyclopentane or cyclohexane, ether such as tetrahydrofran or a mixture of two or more of these solvents or the like.

Alternatively, it may be accomplished by silica gel column chromatography or silica gel thin layer chromatography.

The compound of the formula I''', i.e. the thiazole derivative having cyano group at position 2, is obtained by treating the compound of the formula I'', especially 2-carbamoylthiazole derivative in which R₅ and R₆ are hydrogen atoms, with a dehydrating agent in the usual way. More specifically, the reaction is completed by stirring the compound of the formula I'' and more than equal mole of a dehydrating agent such as p-toluenesul-

fonyl chloride or phosphorus trichloride at 60 to 120°C for about 20 hours. It is preferable to conduct this reaction in the presence of a base such as pyridine.



wherein R₂ and R₃ are as defined above.

Amide such as N,N-dimethylformamide, ether such as tetra-hydrofran or heterocyclic compound having nitrogen atom such as pyridine is employed as a solvent.

Purification of the compound of the formula I''' may be accomplished by recrystallization from halogenated hydrocarbon such as chloroform or lower alcohol such as methanol, ethanol or iso-propanol.

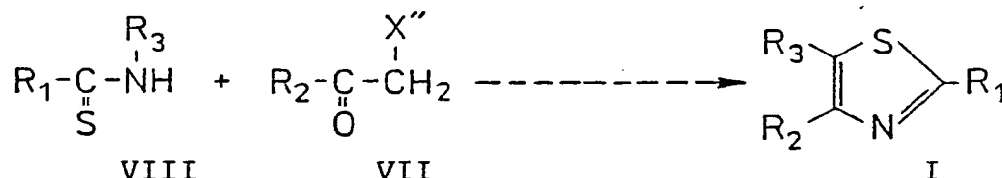
75

Alternatively, it may be accomplished by silica gel column chromatography or thin layer chromatography.

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When the known thioamide of the formula VIII - (in which R₁ and R₃ are as defined above) is reacted with the known halocarbonyl compounds of the formula VII (in which R₂ is as defined above and X'' represents halogen atom) according to the process proposed by Richard H. Willey in Organic Reaction 6 382 (1951), the compound of the formula I may be also obtained.

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wherein R₁, R₂, R₃ and X'' are as defined above.

However, with this known method, the compound of the formula I is obtained in lower yield. This method is unsuitable as industrial process.

The compound of the formula I may be converted into a pharmaceutically acceptable salt by using an appropriate acid.

The appropriate acids which may be used include, for example, inorganic acid such as hydrochloric, hydrobromic, sulfuric, nitric or phosphoric acid, or organic acid such as acetic, pro-

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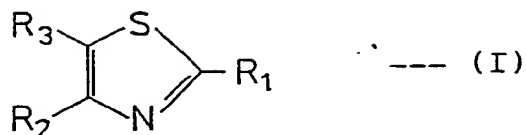
pionic, glycolic, lactic, pyruvic, malonic, succinic, maleic, fumaric, malic, tartaric, citric, benzoic, cinnamic, mandelic, methanesulfonic, hydroxyethanesulfonic, benzenesulfamic, p-toluenesulfamic, cyclohexanesulfamic, salicylic, p-aminosalicylic, 2-phenoxybenzoic or 2-acetoxybenzoic acid.

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The pharmacological effects of the compound of the formula I will now be described;

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wherein R₁, R₂ and R₃ are as defined above.

(1) The contractile force of left atrium was tested according to the method described in Basic Lectures of Medicine Development, Volume V, Pharmacological Test Methods, Part 2, page 535 (1971). A 7-weeks-old male Hartley guinea pig (having a bodyweight of about 350 g) was killed by a blow on a head. The chest was opened and the heart was removed and placed in Krebs-Henseleit solution (prepared by adding distilled water to 6.92 g of sodium chloride, 0.35 g of calcium chloride, 0.29 g of magnesium sulfate, 0.16 g of mono-basic potassium phosphate, 2.1 g

of sodium bicarbonate and 1.8 g of glucose so that the total amount was 1 l) which sufficiently bubbled with oxygen gas. Threads were tied to each tip of the left atrium. One thread was attached to a fixed pin in Magnus's bath and the other to a force-displacement transducer connected to an electric amplifier and recorder. The atrium was stimulated electrically at 0.5 cps, 5 msec and a voltage of 20% above threshold. The preparation is mounted in Krebs-Henseleit solution through which 95% oxygen gas and 5% carbon dioxide was blown. Then the samples were added. The results are shown in Table 1.

Table 1 Effect on Contractile Force
of Isolated Left Atrium

The compound of the formula I	Corresponding Concentration (mol)	Maximum Contractile Force (mg)
R ₁ =-COOCH ₃ , R ₂ =-CH ₃ , R ₃ =4-pyridinyl	1 x 10 ⁻⁴	446 [±] 91
R ₁ =-CONH ₂ , R ₂ =-CH ₃ , R ₃ =4-pyridinyl	1 x 10 ⁻⁴	1,325 [±] 132
R ₁ =-CONHCH ₃ , R ₂ =-CH ₃ , R ₃ =4-pyridinyl	1 x 10 ⁻⁴	510 [±] 45
R ₁ =-CN, R ₂ =-CH ₃ , R ₃ =4-pyridinyl	1 x 10 ⁻⁴	558 [±] 123

(2) The acute toxicity of the compound of the formula I was determined according to the Litchfield-Wilcoxon method, J. Pharm. Exp. Ther., 96, 99 (1949) using 6-weeks-old

male ddY mice (having a bodyweight of 19-24 g) while administrating the sample compound in intraperitoneal injection. The results are shown in Table 2.

Table 2 Acute Toxicity Test Result

The compound of the formula I	LD ₅₀ (mg/kg)
R ₁ =-CONH ₂ , R ₂ =-CH ₃ , R ₃ =4-pyridinyl	230
<p>From the above test results, it was confirmed that the contractile force of isolated left atrium was significantly increased by administration of the compound of the formula I, and the acute toxicity of the compound of the formula I is low.</p>	<p>Electrical amplifier: Model AP-600G supplied by Nippon Kodon Kabushiki Kaisha</p>
<p>The compound of the formula I may be administered to human body orally, by injection - (intravenously, subcutaneously or intramuscularly) or in any other manner. When the compound of the formula I is in the form of solid preparations for oral administration, the preparations may be tablets, granules, powders, capsules or the like. The preparations may contain additives, for example, an excipient such as a saccharide or cellulose preparation, a binder such as starch paste or methyl cellulose, a filler, a disintegrator and so on, all being ones usually used in manufacture of medical preparations.</p>	<p>Programmed electrospphygmometer: supplied by Nippon Kodon Kabushiki Kaisha</p>
<p>In case the compound of the formula I is employed as oral liquid preparations, they may be of any form selected from aqueous preparations for internal use, suspensions, emulsions, syrups, etc., and further they may be in the form of dried products which are dissolved prior to the use.</p>	<p>The invention will be understood more readily with reference to the following examples; however these examples are intended to illustrate the invention and are not to be constituted to limit the scope of the invention. In the examples, the measurements were carried out by using the following apparatuses.</p>
<p>The compound of the formula I may be injected in the form of aqueous solutions, suspensions or oily or queous emulsions, but usually the injections are prepared by dissolving or suspending them in aqueous liquid media such as sterile water of physiological saline solutions.</p>	<p>Melting point: Model MP-1 supplied by Yamato Kagaku Kabushiki Kaisha</p>
<p>If necessary, conventionally used dissolving agents, stabilizers, preservatives, additives for preparing isotonic solutions, etc. may be added to the injections.</p>	<p>Mass analysis: Model M-60 supplied by Kabushiki Kaisha Hitachi Seisakusho</p>
<p>Further, the above-mentioned tests were carried out by using following apparatuses.</p>	<p>Infrared absorption spectrum (IR): Model 260-10 supplied by Kabushiki Kaisha Hitachi Seisakusho</p>
<p>Magnus's bath: supplied by Kabushiki Kaisha Natsume Seisakusho</p>	<p>Nuclear magnetic resonance (NMR): Model FX-270 supplied by Nippon Denshi Kabushiki Kaisha</p>
<p>Recorder: Model WI-680G supplied by Nippon Kodon Kabushiki Kaisha</p>	<p>Elementary analysis: Model MT-2 supplied by Kabushiki Kaisha Yanagimoto Seisakusho</p>
<p>Force displacement transducer: supplied by Nippon Kodon Kabushiki Kaisha</p>	<p>Example 1</p>
	<p>Methyl 4-methyl-5-(4-pyridinyl)-thiazole-2-carboxylate</p>
	<p>(i) Preparation of intermediate, 5-methyl-6-[1-(2,2,2-trichloroethoxycarbonyl)-1,4-dihydro-4-pyridinyl]-2H-1,4-thiazin-3(4H)-one</p>
	<p>2,2,2-Trichloroethylchloroformate (64 ml) was added dropwise to a stirred suspension of 5-methyl-2H-1,4-thiazin-3(4H)-one (50 g) in a mixture of acetonitrile (500 ml) -pyridine (75 ml) under ice-cooling. The mixture was stirred at room temperature for 1 hour and then, poured into cold water - (1.5 l) The resulting precipitates were collected by</p>

filtration, recrystallized from ethanol to give 5-methyl-6-[1-(2,2,2-trichloroethoxycarbonyl)-1,4-dihydro-4-pyridinyl]-2H-1,4-thiazin-3(4H)-one (120 g, yield 80.7%) as pale yellow prisms.

Melting point: 158-160°C

Elementary analysis values as: $C_{13}H_{13}N_2O_2SCl_3$;

Calculated: C = 40.69 H = 3.41 N = 7.29 (%)

Found: C = 40.62 H = 3.37 N = 7.02 (%)

Mass spectrum: M^+ 382

NMR spectrum ($CDCl_3$, TMS) δ :

5 1.986 (3H, s), 3.229 (2H, s),
4.161 (1H, m), 4.800 (4H, m),
6.970 (2H, d), 7.264 (1H, b)

10

IR spectrum

$\nu_{\max}^{KBr} (cm^{-1})$:

: 3200, 3100, 1720, 1670, 1630

(ii) Preparation of intermediate, 5-methyl-6-(4-pyridinyl)-2H-1,4-thiazin-3(4H)-one

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lized from isopropanol to give 5-methyl-6-(4-pyridinyl)-2H-1,4-thiazin-3(4H)-one (0.88 g, yield 76.5%) as pale yellow plates.

Melting point: 187-188.5°C (decomposition)

(A) The mixture of 5-methyl-6-[1-(2,2,2-trichloroethoxycarbonyl)-1,4-dihydro-4-pyridinyl]-2H-1,4-thiazin-3(4H)-one (2.14 g) and sulfur (10.7 g) was stirred at 140°C for 1.5 hours and then cooled to room temperature. The obtained solid was extracted with methanol by using Soxhlet extractor. Methanol was evaporated to dryness, and the residue was dissolved in 50 ml of 2N hydrochloric acid. The insoluble solid was removed by filtration and the filtrate was adjusted to pH 7.2 by addition of 2N sodium hydroxide aqueous solution. The precipitates were collected by filtration and the filtrate was extracted with chloroform (20 ml x 5 times) and was evaporated to dryness. The combined precipitates and the residue were recrystal-

25

Elementary analysis values as: $C_{16}H_{16}N_2OS$;

Calculated: C = 58.22 H = 4.88 N = 13.58 (%)

Found: C = 58.48 H = 4.99 N = 13.53 (%)

Mass spectrum: M^+ 206

NMR spectrum ($CDCl_3$, TMS) δ :

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2.05 (3H, s), 3.43 (2H, s),

35

7.28 (2H, d), 8.61 (2H, d),

35

8.70 (1H, s)

40

IR spectrum

$\nu_{\max}^{KBr} (cm^{-1})$:

: 3200, 3050, 1680, 1580

(B) Zinc powder (1 g) was added to a stirred solution of 5-methyl-6-[1-(2,2,2-trichloroethoxy-carbonyl)-1,4-dihydro-4-pyridinyl]-2H-1,4-thiazin-3(4H)-one (1 g) in formic acid (14 ml), and the reaction mixture was stirred at room temperature for 3 hours. The insoluble solid was removed by filtra-

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tion. The filtrate was evaporated to dryness and the residue was dissolved in water (30 ml). The solution was adjusted to pH 7.0 by addition of 1N sodium hydroxide aqueous solution. The precipitates were extracted with chloroform (although the solution was converted into an emulsion at this extracting step, the operation was facilitated by using a filter aid such as "Avicel"). The extract was

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dried over anhydrous magnesium sulfate and was evaporated under reduced pressure. The crude product was purified by the preparative thin layer chromatography [Merck TLC plate, silica gel 60F₂₅₄t (particles average porous diameter 60A, Fluorescent substance Zn₂SiO₄/Mn), 20x20 cm, t=1 mm, chloroform/methanol = 20/1] to give 200 mg of 5-methyl-6-(4-pyridinyl)-2H-1,4-thiazin-3-(4H)-one. The physical properties were as described above.

(iii) Preparation of intermediate, 2,2-dimethoxy-5-methyl-6-(4-pyridinyl)-2H-1,4-thiazin-3-(4H)-one

m-Chloroperbenzoic acid (5.4 g) was added to a stirred suspension of 5-methyl-6-(4-pyridinyl)-2H-1,4-thiazin-3-(4H)-one (4.5 g) in methanol (180 ml) under ice-cooling, and the mixture was stirred at room temperature for 3 days. The mixture was evaporated to dryness and the residue was extracted with ethyl acetate, and was washed with sodium bicarbonate aqueous solution and water, dried over magnesium sulfate and concentrated under reduced pressure. The residue was chromatographed on activated alumina column (100 g, 300 mesh, manufactured by Wako Junyaku Kabushiki Kaisha) and eluted with chloroform (30 ml). The eluate was concentrated under reduced pressure and the residue was washed with ether (100 ml), and was collected by filtration. m-Chloroperbenzoic acid (2.8 g) was added to a stirred solution of the resulting precipitates in methanol (100 ml) at room temperature, and the reaction mixture was stirred at room temperature for 2 hours. The reaction mixture was evaporated to dryness and the residue was dissolved in ethyl acetate. The extract was washed with sodium bicarbonate aqueous solution and successively with water, dried over magnesium sulfate and concentrated under reduced pressure. The residue was washed with ether, and the resulting powder was collected by filtration to give 2,2-dimethoxy-5-methyl-6-(4-pyridinyl)-2H-1,4-thiazin-(4H)-one (2 g, yield 35.0%) as pale yellow powder.

Melting point: 154-155°C

NMR spectrum (CDCl₃, TMS) δ :

5 2.03 (3H, s), 3.59 (6H, s),
7.25 (2H, dd), 8.43 (1H, s),
8.62 (2H, dx2)

10

(iv) Preparation of methyl 4-methyl-5-(4-pyridinyl)-thiazole-2-carboxylate

2,2-Dimethoxy-5-methyl-6-(4-pyridinyl)-2H-1,4-thiazin-3-(4H)-one (200 mg) was dissolved in 2N hydrochloric acid (2 ml), and the mixture was allowed to stand at room temperature for 30 minutes. Then, the solution was made alkaline by addition of 2N aqueous sodium hydroxide under ice-cooling, and the resulting precipitates were collected by filtration. The precipitates were washed with water, air-dried, and was recrystallized from benzene-petroleum ether to give methyl 4-methyl-5-(4-pyridinyl)-thiazole-2-carboxylate (158 mg, yield 90%) as colorless needles.

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Melting point: 121-123°C

Elementary analysis values as: C₁₁H₁₀N₂O₂S;

30 Calculated: C = 56.39 H = 4.30 N = 11.95 (%)

Found: C = 56.33 H = 4.29 N = 11.62 (%)

Mass spectrum: M⁺ 234

35

NMR spectrum (CDCl₃, TMS) δ :

2.65 (3H, s), 4.04 (3H, s),
40 7.40 (2H, dd), 8.72 (2H, dx2),

IR spectrum

45

$\nu_{\text{max}}^{\text{KBr}} (\text{cm}^{-1})$:

50

: 1750, 1720, 1600

In the same manner as described in Example 1 (i)-(iv), the compounds of Example 2 to 4 are obtained.

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Example 2

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Methyl 4-methyl-5-(3-methyl-4-pyridinyl)-thiazole-2-carboxylate

Melting point: 95.5-96.3°C

5-Methyl-2H-1,4-thiazine-3(4H)-one was treated with 3-methyl-pyridine to give methyl 4-methyl-5-(3-methyl-4-pyridinyl)-thiazole-2-carboxylate as colorless prisms.

NMR spectrum (CDCl₃, TMS) δ :

5 2.24 (3H, s), 2.37 (3H, s),
4.04 (3H, s), 7.19 (1H, d),
8.54 (1H, d), 8.60 (1H, s)

10

IR spectrum

$\nu_{\text{max}}^{\text{KBr}} (\text{cm}^{-1}) :$

: 1735

Melting point: 193-194°C

Example 3

20

NMR spectrum (CDCl₃, TMS) δ :

Methyl 4-methyl-5-(3-cyano-4-pyridinyl)-thiazole-2-carboxylate

25

2.55 (3H, s), 4.05 (3H, s),
7.47 (1H, d), 8.90 (1H, d),

5-Methyl-2H-1,4-thiazine-3(4H)-one was treated with 3-cyano-pyridine to give methyl 4-methyl-5-(3-cyano-4-pyridinyl)-thiazole-2-carboxylate as colorless columns.

9.03 (1H, s)

IR spectrum

30

$\nu_{\text{max}}^{\text{KBr}} (\text{cm}^{-1}) :$

35

: 2230, 1710

Melting point: 116-117°C

Example 4

40

NMR spectrum (CDCl₃, TMS) δ :

Methyl 4-methyl-5-(4-quinolinyl)-thiazole-2-carboxylate

45

2.37 (3H, s), 4.07 (3H, s),
7.41 (1H, d), 7.60-8.23 (4H, m),

5-Methyl-2H-1,4-thiazine-3(4H)-one was treated with quinoline to give methyl 4-methyl-5-(4-quinolinyl)-thiazole-2-carboxylate as colorless granules.

9.01 (1H, d)

IR spectrum

50

$\nu_{\text{max}}^{\text{KBr}} (\text{cm}^{-1}) :$

55

: 1730, 1710

Example 5

4-Methyl-5-(4-pyridinyl)-thiazole-2-carboxamide

Methyl 4-methyl-5-(4-pyridinyl)-thiazole-2-carboxylate (3 g) was added to a stirred ammonia- (28%, 85 ml) at room temperature, and the mixture was stirred at room temperature for overnight. The resulting precipitate was collected by filtration,

washed with water, and air dried. The precipitate were recrystallized from ethanol to give 4-methyl-5-(4-pyridinyl)-thiazole-2-carboxamide (2.3 g, yield 82%) as pale yellow needles.

Melting point: 222.5-224°C

Elementary analysis values as: $C_{10}H_{11}N_3OS$;

Calculated: C = 54.77 H = 4.13 N = 19.16 (%)

Found: C = 54.77 H = 4.11 N = 18.73 (%)

Mass spectrum: M^+219

IR spectrum

$\nu_{\max}^{KBr} (cm^{-1})$:

: 3450, 3200, 1670, 1580

Example 6

N,4-Dimethyl-5-(4-pyridinyl)-thiazole-2-carboxamide

Methyl amine (0.5 ml) was added to a stirred solution of methyl 4-methyl-5-(4-pyridinyl)-thiazole-2-carboxylate (250 mg) in absolute ethanol (10 ml) at room temperature, and the mixture was stirred at room temperature for overnight. The mixture was evaporated to dryness and the residue was recrystallized from iso-propanol to give N,4-dimethyl-5-(4-pyridinyl)-thiazole-2-carboxamide (200 mg, yield 80.0%) as pale yellow needles.

Melting point: 184-186°C

Elementary analysis values as: $C_{11}H_{11}N_3OS$;

Calculated: C = 56.63 H = 4.75 N = 18.01 (%)

Found: C = 56.69 H = 4.80 N = 17.50 (%)

NMR spectrum ($CDCl_3$, TMS) δ :

2.56 (3H, s), 3.04 (3H, d),

7.26 (1H, b), 7.40 (2H, dd),

8.70 (2H, dx2),

IR spectrum

$\nu_{\max}^{KBr} (cm^{-1})$:

: 3200, 3100, 1670, 1600

Example 7

4-Methyl-5-(4-pyridinyl)-thiazole-2-carboxanilide

n-Butyllithium (14-17% in hexane, 1.2 ml) was added to dropwise a stirred solution of aniline (0.12 ml) in dry tetrahydrofuran (3 ml) at room temperature under nitrogen atmosphere. Then, methyl 4-

methyl-5-(4-pyridinyl)-thiazole-2-carboxylate (200 mg) was added to a stirred solution at room temperature under nitrogen atmosphere, and the mixture was stirred in the same condition for 1 hour. The resulting precipitates were collected by filtration, and the precipitates were washed with water, and air-dried. The precipitate was recrystallized from benzenepetroleum ether to give 4-methyl-5-(4-pyridinyl)-thiazole-2-carboxanilide (80 mg, yield 33.3%) as colorless plates.

Melting point: 196-197°C

Elementary analysis values as: $C_{16}H_{13}N_2OS$;

Calculated: C = 65.06 H = 4.43 N = 14.22 (%)

Found: C = 64.98 H = 4.43 N = 13.82 (%)

NMR spectrum ($CDCl_3$, TMS) δ :

2.62 (3H, s), 7.18-7.43 (5H, m),

7.75 (2H, dd), 8.72 (2H, dx2),

9.05 (1H, s)

IR spectrum

10

$\nu_{\max}^{KBr} (cm^{-1})$:

15

: 3220, 3100, 1660, 1590

Example 8

4-Methyl-N-(2-dimethylaminoethyl)-5-(4-pyridinyl)-thiazole-2-carboxamide hydrochloride

N,N-Dimethylethylenediamine (1.4 ml) was added to a suspension of methyl 4-methyl-5-(4-pyridinyl)-thiazole-2-carboxylate (200 mg) in water - (2.8 ml) at room temperature, and the mixture was

stirred at room temperature for overnight. The mixture was evaporated to dryness and the residue was chromatographed on an activated alumina column and eluted with chloroform to give 4-methyl-N-(2-dimethylaminoethyl)-5-(4-pyridinyl)-thiazole-2-carboxamide as yellow oil.

IR spectrum

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25

$\nu_{\max}^{KBr} (cm^{-1})$:

: 3200, 1650, 1590, 1530 4-Methyl-N-(2-dimethylaminoethyl)-5-(4-pyridinyl)-thiazole-2-carboxamide was adjusted to pH 1 to 2 by addition of conc. hydrochloric acid, and the solution was evaporated to dryness. The residue was washed with ethanol to give 4-methyl-N-(2-dimethylaminoethyl)-5-(4-pyridinyl)-thiazole-2-carboxamide hydrochloride as colorless powder.

Melting point: over 220°C (decomposition)

Elementary analysis values as $C_{14}H_{20}N_4OSCl_2 \cdot 2H_2O$;

Calculated: C = 43.18 H = 6.21 N = 14.39 (%)

Found: C = 43.62 H = 5.74 N = 14.38 (%)

NMR spectrum ($DMSO-d_6$, TMS) δ :

35

2.67 (3H, s), 2.81 (3H, s),

2.83 (3H, s), 3.30 (2H, m),

3.70 (2H, m), 8.15 (2H, dd)

40

8.95 (2H, dx2), 9.24 (1H, t)

Example 9

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N,N-diethyl-4-methyl-5-(4-pyridinyl)-thiazole-2-carboxamide

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(i) Preparation of 4-methyl-5-(4-pyridinyl)-thiazole-2-carboxylic acid

Potassium hydroxide (1 g) was added to a stirred solution of methyl 4-methyl-5-(4-pyridinyl)-thiazole-2-carboxylate (2 g) in methanol (50 ml) at room temperature and the mixture was stirred at

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room temperature for 1.5 hours, diluted with water - (150 ml), and washed with chloroform. Then the solution was adjusted to pH 3 to 4 by addition of 2N hydrochloric acid aqueous solution. The resulting precipitate was collected by filtration, and air-dried to give 4-methyl-5-(4-pyridinyl)-thiazole-2-carboxylic acid (1.8g, 96%) as colorless powder.

Melting point: 133-134°C (decomposition)

Elementary analysis values as: $C_{10}H_8N_2O_2S$;

5 Calculated: C = 54.33 H = 3.66 N = 12.72 (%)

Found: C = 53.95 H = 3.54 N = 12.37 (%)

IR spectrum

10

$\nu_{\max}^{KBr} (cm^{-1}) :$

15

: 1705, 1630, 1610, 1520

(ii) Preparation of N,N-diethyl-4-methyl-5-(4-pyridinyl)-thiazole-2-carboxamide

with ethyl acetate-n-hexane = 6:4 to give N,N-diethyl-4-methyl-5-(4-pyridinyl)-thiazole-2-carboxamide (60 mg, yield 48%) as colorless needles.

20

Melting point: 87.5-89°C

Elementary analysis values as: $C_{14}H_{17}N_3OS$;

25 Calculated: C = 61.06 H = 6.22 N = 15.26 (%)

Found: C = 60.68 H = 6.25 N = 14.62 (%)

NMR spectrum ($CDCl_3$, TMS) δ :

30

1.3 (6H, t), 2.6 (3H, s),

3.3-4.4 (4H, m), 7.5 (2H, dd),

35

8.8 (2H, dx2)

IR spectrum

$\nu_{\max}^{KBr} (cm^{-1}) :$

: 1610, 1590

Example 10

2',6'-dimethyl-4-methyl-5-(4-pyridinyl)-thiazole-2-carboxanilide

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the mixture, and was stirred at room temperature for overnight. The mixture was diluted with chloroform, washed with water, dried over magnesium sulfate, and evaporated to dryness. The residue was recrystallized from iso-propanol to give 2',6'-dimethyl-4-methyl-5-(4-pyridinyl)-thiazole-2-carboxanilide (180 mg, yield 31%) as colorless plates.

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2,6-Dimethyl aniline hydrochloride (230 mg) was added to a stirred suspension of 4-methyl-5-(4-pyridinyl)-thiazole-2-carbonylchloride hydrochloride (230 mg) in dry toluene (4 ml) at room temperature, and triethylamine (1.8 ml) was added to

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Melting point: 192-194°C

Elementary analysis values as: $C_{15}H_{17}N_3OS$;

Calculated: C = 66.84 H = 5.29 N = 12.99 (%)

Found: C = 66.30 H = 5.30 N = 12.43 (%)

2.33 (6H, s), 2.63 (2H, s),

7.17 (3H, m), 7.42 (2H, dd),

5 8.59 (1H, s), 8.72 (2H, dx2)

IR spectrum

NMR spectrum ($CDCl_3$, TMS) δ :

10

$\nu_{\max}^{KBr} (cm^{-1})$:

15

:3350, 1680, 1590

Example 11

1-[4-methyl-5-(4-pyridinyl)-thiazolyl-2-carbonyl]-piperidine

Piperidine (1.4 ml) was added to a stirred suspension of 4-methyl-5-(4-pyridinyl)-thiazole-2-carbonylchloride hydrochloride (400 mg) in dry toluene (4 ml) at room temperature, and the mixture was stirred at room temperature for overnight. The mixture was treated in the same manner as described in Example 6, and was recrystallized from ethyl acetate-n-hexane to give 1-[4-methyl-5-(4-pyridinyl)-thiazolyl-2-carbonyl]-piperidine (100 mg, yield 24%) as pale yellow granules.

Melting point: 139-142°C (decomposition)

Elementary analysis values as: $C_{15}H_{17}N_3OS$;

Calculated: C = 62.68 H = 5.96 N = 14.62 (%)

Found C = 62.77 H = 6.01 N = 14.09 (%)

NMR spectrum ($CDCl_3$, TMS) δ :

1.70 (6H, m), 2.58 (3H, s),

3.75 (2H, m), 4.25 (2H, m),

7.40 (2H, dd), 8.70 (2H, dx2)

IR spectrum

35

$\nu_{\max}^{KBr} (cm^{-1})$:

40

: 1610, 1590

Example 12

2-cyano-4-methyl-5-(4-pyridinyl)-thiazole

p-Toluenesulfonyl chloride (860 mg) was added to a stirred suspension of 4-methyl-5-(4-pyridinyl)-thiazole-2-carboxamide (450 mg) in dry pyridine (5 ml) under ice-cooling, and the mixture was stirred at 90°C for 21 hours. The mixture was evaporated to dryness, and the residue was dissolved 2N hydrochloric acid aqueous solution (10-15 ml), washed twice with chloroform, and made alkaline by addition of 2N sodium hydroxide aqueous solu-

tion. The resulting precipitates were extracted with chloroform, washed with water, dried over magnesium sulfate, and evaporated to dryness. The residue was washed with petroleum ether, and was recrystallized from iso-propanol (twice) to give 2-cyano-4-methyl-5-(4-pyridinyl)-thiazole. (220 mg, yield 53%) as pale yellow needles.

Melting point: 169-170°C

Elementary analysis values as: $C_{10}H_7N_3S$;

Calculated: C = 59.67 H = 3.50 N = 20.88 (%)

Found: C = 59.36 H = 3.44 N = 20.45 (%)

22

NMR spectrum (CDCl₃, TMS) δ :

2.63 (3H, s), 7.39 (2H, dd),

8.76 (2H, dx2)

5

IR spectrum

$\nu_{\text{max}}^{\text{KBr}} (\text{cm}^{-1})$:

: 2240, 1600

Example 13

Methyl 4-methyl-5-(2-thiazolyl)-thiazole-2-carboxylate

(i) Preparation of intermediate, 2-methoxy-5-methyl-2H-1,4-thiazin-3(4H)-one

m-Chloroperbenzoic acid (28.8 g) was added dropwise to a stirred solution of 5-methyl-2H-1,4-thiazin-3(4H)-one (15 g) in methanol (300 ml) under ice-cooling and the mixture was stirred at room temperature for 1 day. After the mixture was evaporated to dryness, the residue was extracted with ethyl acetate, washed with saturated sodium bicarbonate aqueous solution and dried over anhydrous magnesium sulfate. The solvent was evaporated to dryness to give 2-methoxy-5-methyl-2H-1,4-thiazin-3(4H)-one (11.3g, yield 61%) as pale yellow powder.

Melting point: 157-160°C

(ii) Preparation of intermediate, 2,2-Dimethoxy-5-methyl-2H-1,4-thiazin-3(4H)-one

m-Chloroperbenzoic acid (25.6 g) was added dropwise to a stirred solution of 2-methoxy-5-methyl-2H-1,4-thiazin-3(4H)-one (16 g) in methanol (500 ml) under ice-cooling and the mixture was stirred at room temperature for 4 hours. Then, the reaction mixture was made alkaline by addition of anhydrous potassium carbonate and evaporated to dryness. The residue was extracted twice with chloroform and the combined extracts were washed with water, dried over anhydrous magne-

sium sulfate and evaporated to dryness. The residue was triturated with ethyl ether to give 2,2-dimethoxy-5-methyl-2H-1,4-thiazin-3(4H)-one (10 g, yield 53%) as pale yellow plates.

Melting point: 86-88°C

(iii) Preparation of intermediate, 2,2-dimethoxy-5-methyl-6-(3-ethoxycarbonyl-2,3-dihydro-2-thiazolyl)-2H-1,4-thiazin-3(4H)-one

Ethyl chloroformate (1.0 g) was added dropwise to a stirred solution of thiazole (0.79 g) in dichloromethane (10 ml) under ice-cooling and the mixture was stirred at room temperature for 0.5 hour. Then, 2,2-dimethoxy-5-methyl-2H-1,4-thiazin-3(4H)-one (1.47 g) was added, and the mixture was stirred at room temperature for 3 hours. The solution was washed with saturated sodium bicarbonate aqueous solution, dried over magnesium sulfate and evaporated to dryness. The residue was chromatographed on silica gel (Merck Lobar column size B: Art.10401, solvent: ethyl acetate-n-hexane = 2:3) to give 2,2-dimethoxy-5-methyl-6-(3-ethoxycarbonyl-2,3-dihydro-2-thiazolyl)-2H-1,4-thiazin-(4H)-one (510 mg, yield 19%) as colorless prisms.

Melting point: 148-149°C

(iv) Preparation of intermediate, 2,2-dimethoxy-5-methyl-6-(2-thiazolyl)-2H-1,4-thiazine-3-one

DDQ (206.4 mg) was added dropwise to a stirred solution of 2,2-dimethoxy-5-methyl-6-(3-ethoxycarbonyl-2,3-dihydro-2-thiazolyl)-2H-1,4-thiazine-3(4H)-one (300 mg) in dichloromethane (5 ml) at room temperature, and the mixture was stirred at room temperature for 1 hour. Then, the insoluble materials were filtered off and the solvent was evaporated to dryness.

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The residue was chromatographed on silica gel (Merck Lobar column size B: Art.10401, solvent: ethyl acetate-n-hexane = 2:3) to give 2,2-dimethoxy-5-methyl-6-(2-thiazolyl)-2H-1,4-thiazin-3-(4H)-one (144 mg, yield 61%) as colorless column.

Melting point: 127-128°C

(v) Methyl 4-methyl-5-(2-thiazolyl)-thiazole-2-carboxylate

A mixture of 2,2-dimethoxy-5-methyl-6-(2-thiazolyl)-2H-1,4-thiazin-3-(4H)-one (80 mg) and 2N hydrochloric acid solution (2 ml) was stirred at room temperature for 6 hours. Then the mixture was neutralized with saturated sodium bicarbonated aqueous solution and the resulting precipitates were collected to give methyl 4-methyl-5-(2-thiazolyl)-thiazole-2-carboxylate (58 mg, yield 82.2%) as colorless powder.

Melting point: 129-129.5°C

As is apparent from the foregoing description, the thiazole derivative of the present invention is a compound not described in any literature, the contractile force of isolated left atrium was significantly increased by administration of the thiazole deriva-

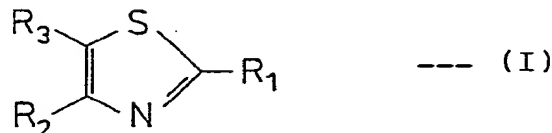
tive of the present invention, and the acute toxicity of the thiazole derivative of the present invention is low. Accordingly, the thiazole derivative of the present invention is effective in curing and preventing heart diseases, especially cardiac insufficiency.

As for the process of the present invention, it is advantageous from the industrial view point since the thiazole derivative of the present invention can be prepared from relatively easily available starting compounds in high yield by a relatively easy operation.

While the preferred form of the present invention has been described, it is to be understood that modifications will be apparent to those skilled in the art without departing from the true spirit of the invention. The scope of the invention, therefore, is to be determined solely by the following claims.

Claims

1. A thiazole derivative represented by the following formula I and pharmaceutically acceptable acid addition salt thereof:



wherein R_1 represents $-\text{COOR}_4$, $-\text{CON} \begin{array}{c} R_5 \\ \diagup \\ R_6 \end{array}$ or cyano

wherein R_4 represents hydrogen or lower alkyl, and

R_5 and R_6 may be same or different and represent hydrogen, lower alkyl, aryl, amino-lower alkyl, N-lower alkylamino-lower alkyl, N,N-di-lower alkylamino-lower alkyl or R_5 and R_6 are joined to form N-containing heterocyclic group;

R_2 represents hydrogen or lower alkyl; and

R_3 represents N-containing heterocyclic group.

2. A compound of Claim 1, wherein R_1 is carboxy, methoxycarbonyl, carbamoyl, methylaminocarbonyl, phenylaminocarbonyl, diethylaminocarbonyl or cyano.

3. A compound of Claim 1, wherein R_2 is methyl.

4. A compound of Claim 1, wherein R_3 is 4-

pyridinyl, 3-methyl-4-pyridinyl, 3-cyano-4-pyridinyl, quinolinyl or thiazolyl.

5. 4-Methyl-5-(4-pyridinyl)-thiazole-2-carboxylic acid, according to Claim 1.

6. Methyl 4-methyl-5-(4-pyridinyl)-thiazole-2-carboxylate, according to Claim 1.

7. Methyl 4-methyl-5-(3-methyl-4-pyridinyl)-thiazole-2-carboxylate, according to Claim 1.

8. Methyl 4-methyl-5-(3-cyano-4-pyridinyl)-thiazole-2-carboxylate, according to Claim 1.

9. Methyl 4-methyl-5-(4-quinolinyl)-thiazole-2-carboxylate, according to Claim 1.

10. 4-Methyl-5-(4-pyridinyl)-thiazole-2-carboxamide, according to Claim 1.

11. N,4-Dimethyl-5-(4-pyridinyl)-thiazole-2-carboxamide, according to Claim 1.

12. 4-Methyl-5-(4-pyridinyl)-thiazole-2-carboxanilide, according to Claim 1.

13. 4-Methyl-N-(2-dimethylaminoethyl)-5-(4-pyridinyl)-thiazole-2-carboxamide, according to Claim 1.

14. N,N-Diethyl-4-methyl-5-(4-pyridinyl)-thiazole-2-carboxamide, according to Claim 1.

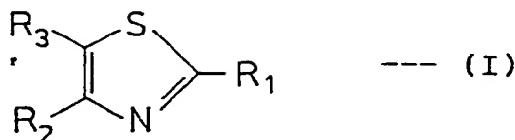
15. 2',6'-Dimethyl-4-methyl-5-(4-pyridinyl)-thiazole-2-carboxamide, according to Claim 1.

16. 1-[4-Methyl-5-(4-pyridinyl)-thiazolyl-2-carbonyl]-piperidine, according to Claim 1.

17. 2-Cyano-4-methyl-5-(4-pyridinyl)-thiazole, according to claim 1.

18. Methyl 4-methyl-5-(2-thiazolyl)-2-carboxylate, according to Claim 1.

19. A cardiotonic agent comprising a pharmaceutically acceptable excipient and, as an active component thereof, an effective amount of thiazole derivative having the following formula I:



wherein R_1 represents $-COOR_4$, $-CON(R_5)(R_6)$ or cyano

wherein R_4 represents hydrogen or lower alkyl, and

R_5 and R_6 may be same or different and represent hydrogen, lower alkyl, aryl, amino-lower alkyl, N-lower alkylamino-lower alkyl, N,N-di-lower alkylamino-lower alkyl or R_5 and R_6 are joined to form N-containing heterocyclic group;

R_2 represents hydrogen or lower alkyl; and

R_3 represents N-containing heterocyclic group.

20. An agent according to Claim 19, wherein the active component is 4-methyl-5-(4-pyridinyl)-thiazole-2-carboxylic acid.

21. An agent according to Claim 19, wherein the active component is methyl 4-methyl-5-(4-pyridinyl)-thiazole-2-carboxylate.

22. An agent according to Claim 19, wherein the active component is methyl 4-methyl-5-(3-methyl-4-pyridinyl)-thiazole-2-carboxylate.

23. An agent according to Claim 19, wherein the active component is methyl 4-methyl-5-(3-cyano-4-pyridinyl)-thiazole-2-carboxylate.

24. An agent according to Claim 19, wherein the active component is methyl 4-methyl-5-(2-

thiazolyl)-thiazole-2-carboxylate.

25. An agent according to Claim 19, wherein the active component is 4-methyl-5-(4-pyridinyl)-thiazole-2-carboxamide.

26. An agent according to Claim 19, wherein the active component is N,4-dimethyl-5-(4-pyridinyl)-thiazole-2-carboxamide.

27. An agent according to Claim 19, wherein the active component is 4-methyl-5-(4-pyridinyl)-thiazole-2-carboxanilide.

28. An agent according to Claim 19, wherein the active component is 4-methyl-N-(2-dimethylaminoethyl)-5-(4-pyridinyl)-thiazole-2-carboxamide.

29. An agent according to Claim 19, wherein the active component is N,N-diethyl-4-methyl-5-(4-pyridinyl)-thiazole-2-carboxamide.

30. An agent according to Claim 19, wherein the active component is 2',6'-dimethyl-4-methyl-5-(4-pyridinyl)-thiazole-2-carboxamide.

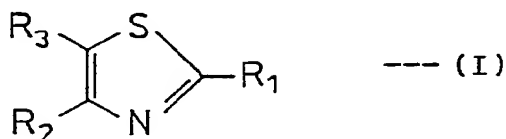
31. An agent according to Claim 19, wherein the active component is 1-[4-methyl-5-(4-pyridinyl)-thiazolyl-2-carbonyl]-piperidine.

32. An agent according to Claim 19, wherein the active component is 2-cyano-4-methyl-5-(4-pyridinyl)-thiazole.

33. An agent according to Claim 19, wherein the

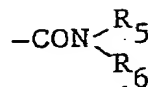
active component is methyl 4-methyl-5-(2-thiazolyl)-thiazole-2-carboxylate.

34. A process for preparation of a thiazole derivative represented by the following formula I



wherein R₁ represents -COOR₄,

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or cyano wherein R₄ represents hydrogen or lower alkyl, and

R₅ and R₆ may be same or different and represent hydrogen, lower alkyl, aryl, amino-lower alkyl, N-lower alkylamino-lower alkyl, N,N-di-lower alkylamino-lower alkyl or R₅ and R₆ are joined to

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form N-containing heterocyclic group;

R₂ represents hydrogen or lower alkyl; and R₃ represents N-containing heterocyclic group,

which comprises the steps of reacting a thiazine derivative of the formula VI

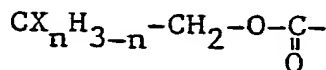
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--- (VI)

wherein R₂ is as defined above, with a compound of the formula A-X' wherein A represents a group

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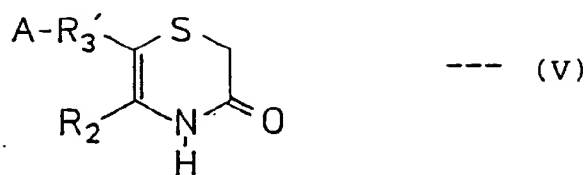
in which X represents a halogen atom and n is number of 0 to 3 and X' represents a halogen atom which may be the same as or different from X, and

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a compound of the formula R₃-H wherein R₃ is as defined above, to produce a compound of the formula V

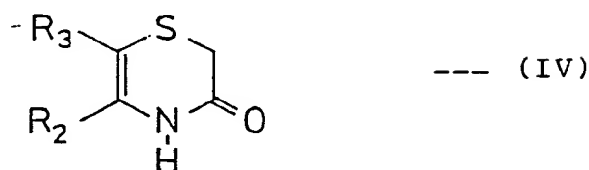
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wherein R_2 and A are as defined above and R_3' represents dihydroform residue of N-containing heterocyclic group; aromarizing the compound of the formula V to produce a compound of the formula IV

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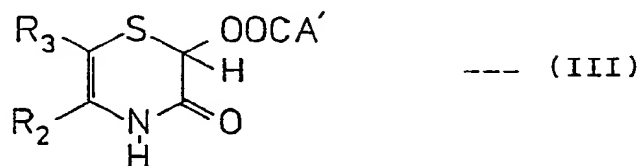


wherein R_2 and R_3 are as defined above;

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reacting the compound of the formula IV with a peroxycarboxylic acid represented by the formula

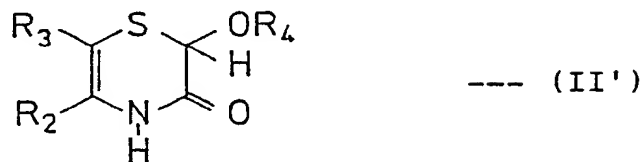
$\text{A}'\text{-COOOH}$ wherein A' is lower alkyl, alicyclic residue or aryl to produce a compound of the formula III



wherein R_2 , R_3 and A' are as defined above;

reacting the compound of the formula III with a nucleophilic reagent represented by the formula $\text{R}_4\text{-OH}$ wherein R_4 is as defined above, to produce a compound of the formula II'

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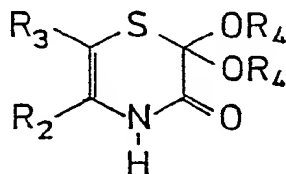


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wherein R_2 , R_3 and R_4 are as defined above;

reacting the compound of the formula II' with a compound of the formula $A'-COOOH$ wherein A' is



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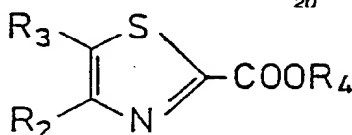
as defined above and a compound of the formula R_4-OH wherein R_4 is as defined above, to produce a compound of the formula II

--- (II)

wherein R_2 , R_3 and R_4 are as defined above;

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reacting the compound of the formula II with an acid, to produce a compound of the formula I'



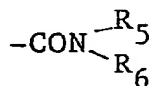
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--- (I')

wherein, R_2 , R_3 and R_4 are as defined above;

optionally the $-COOR_4$ group of the compound of the formula I' may be converted into

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group (in which R_5 and R_6 as defined above) or cyano by methods known in the art.

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35. A process according to Claim 34, wherein a thiazine derivative of the formula VI is 5-methyl-2H-1,4-thiazine-3(4H)-one.

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36. A process according to Claim 34, wherein a compound of the formula $A-X'$ is 2,2,2-trichloroethylchloroformate.

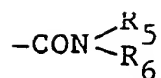
37. A process according to Claim 34, wherein a compound of the formula R_2-H is pyridine, 3-methylpyridine, 3-cyanopyridine, quinoline or thiazole.

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41. A process according to Claim 34, wherein the acid is hydrochloric acid.

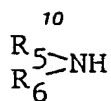
42. A process according to Claim 34, wherein conversion of the

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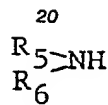
group is effected by reacting the compound of the formula



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with the compound of the formula I'.

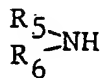
43. A process according to Claim 34, wherein the compound of the formula



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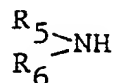
is ammonia, methyl amine, aniline, 2,6-dimethylaniline, N,N-dimethylaminoethylamine, diethylamine or piperidine.

44. A process according to Claim 34, wherein conversion of the cyano is effected by reacting the compound of the formula



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45. A process according to Claim 34, wherein the compound of the formula



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is ammonia.

46. A process according to Claim 34, wherein a dehydrating agent is p-toluenesulfonylchloride.

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